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## CLAIM AMENDMENTS

## Claims 1 through 18 (canceled)

- (Currently amended) An isolated nucleic acid 1 sequence from the ATI region of modified vaccinia Ankara virus that 2 includes at least one restriction enzyme recognition site as an 3 insertion site for a heterologous sequence and that hybridizes under stringent conditions to the nucleic acid sequence of SEQ ID 5 NO:1 or its complementary strand which includes multiple cloning sites inserted into an open reading frame or the ECORI site of the 7 isolated nucleic acid sequence, said nucleic acid sequence capable 8 of integration of [[the]] a heterologous sequence through 9 homologous recombination into an open reading frame or an ECORI 10 site of the ATI region of an orthopoxvirus without interfering with 11 its viral propagation or replication efficiency. 12
  - 20. (Currently amended) The nucleic acid sequence defined in claim 19 that includes as the insertion site for the multiple cloning sites an ECORI site corresponding to position 1063 of SEQ ID NO:1 that hybridizes under stringent conditions to the nucleic acid sequence of SEQ ID NO:1 or its complementary strand.
- 21. (Currently amended) An isolated fragment of a nucleic acid sequence from the ATI region of modified vaccinia

  Ankara virus consisting essentially of at least 200 base pairs of

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replication efficiency.

the nucleic acid sequence that is SEQ ID NO:1, that is at least 70% homologous to SEQ ID NO:1 and that includes at least one 5 restriction enzyme recognition site as an insertion site for a 6 heterologous sequence, which includes multiple cloning sites 7 inserted into an open reading frame or the ECORI site of the 8 isolated fragment of a nucleic acid sequence, said isolated 9 fragment of a nucleic acid sequence capable of integration of 10 [[the]] a heterologous sequence through homologous recombination 11 into an open reading frame or an ECORI site of the ATI region of an 12

orthopoxvirus without interfering with its viral propagation or

- 22. (Currently amended) The isolated fragment of a nucleic acid sequence defined in claim 21 that includes as the insertion site for the multiple cloning sites an ECORI site corresponding to position 1063 of SEQ ID NO:1.
  - heterologous sequence into an open reading frame of or into the ECORI site of the ATI region of an orthopoxviral genome having an ATI region, said vector including an isolated nucleic acid sequence from the ATI region of modified vaccinia Ankara virus that includes at least one restriction enzyme recognition site as an insertion site for a heterologous nucleic acid sequence, that hybridizes under stringent conditions to the nucleic acid sequence of SEQ ID NO:1 or its complementary strand, which includes multiple cloning

- sites inserted into an open reading frame or the ECORI site of the
  isolated nucleic acid sequence, and that is said nucleic acid
  sequence and that is capable of integration of the heterologous
  sequence through homologous recombination into an open reading
  frame or the ECORI site of the ATI region of an orthopoxvirus
  without interfering with its viral propagation or replication
  efficiency.
- 24. (Previously presented) The vector defined in claim
  23 wherein additionally at least one transcriptional control
  3 element is included in the insertion site.
- 25. (Previously presented) The vector defined in claim
  2 23 wherein the insertion site is the restriction site ECOR1.
- 26. (Previously presented) The vector defined in claim
  24 wherein the at least one transcriptional control element is
  3 obtained from a poxvirus genome or is a consensus sequence from a
  4 poxvirus genome.
- further comprising at least one heterologous sequence inserted
  within the insertion site into an open reading frame or the ECORI
  site of the isolated nucleic acid sequence, said heterologous
  nucleic acid sequence functionally associated with a
  transcriptional control element thereof.

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- 28. (Previously presented) The vector defined in claim
  27 wherein the heterologous nucleic sequence is selected from the
  3 group consisting of marker genes, therapeutic genes, host range
  4 genes and genes encoding immunogenic epitopes.
- 29. (Previously presented) The vector defined in claim
  27 comprising a recombinogenic sequence, which flanks one or more
  3 heterologous sequences encoding marker genes, host range genes,
  4 and/or a transcriptional element thereof.
  - heterologous sequence into an open reading frame of or into the ECORI site of the ATI region of an orthopoxviral genome having an ATI region, said vector including an isolated fragment of a nucleic acid sequence from the ATI region of modified vaccinia Ankara virus consisting essentially of at least 200 base pairs of the nucleic acid sequence that is SEQ ID NO:1, that is at least 70% homologous to SEQ ID NO:1 and that includes at least one restriction enzyme recognition site as an insertion site for the heterologous sequence, which includes multiple cloning sites inserted into an open reading frame or the ECORI site of the isolated fragment of a nucleic acid sequence, said isolated fragment of the nucleic acid sequence capable of integration of the heterologous sequence through homologous recombination into an open reading frame or an

- ECORI site of the ATI region of an orthopoxvirus without
- interfering with its viral propagation or replication efficiency.
- 1 31. (Previously presented) The vector defined in claim
- 30 wherein additionally at least one transcriptional control
- element is included in the insertion site.
- 32. (Previously presented) The vector defined in claim
- 30 wherein the insertion site is the restriction site ECOR1.
- 1 33. (Previously presented) The vector defined in claim
- 31 wherein the at least one transcriptional control element is
- obtained from a poxvirus genome or is a consensus sequence from a
- 4 poxvirus genome.
- 1 34. (Currently amended) The vector defined in claim 30
- further comprising at least one heterologous sequence inserted
- 3 within the insertion site into an open reading frame or the ECORI
- site of the isolated nucleic acid sequence, said heterologous
- 5 nucleic acid sequence functionally associated with a
- transcriptional control element thereof.
- 35. (Previously presented) The vector defined in claim
- 34 wherein the heterologous sequence is selected from the group
- 3 consisting of marker genes, therapeutic genes, host range genes and
- genes encoding immunogenic epitopes.

- 36. (Previously presented) The vector defined in claim
  34 comprising a recombinogenic nucleic acid sequence, which flanks
  one or more heterologous sequences encoding marker genes, host
  range genes, and/or a transcriptional element thereof.
- 1 37. (Currently amended) A recombinant orthopoxvirus

  2 having an ATI region, comprising in an open reading frame of or in

  3 the ECORI site of its ATI region an integrated heterologous nucleic

  4 acid sequence wherein said integrated heterologous nucleic acid

  5 sequence does not interfere with viral propagation and/or

  6 replication efficiency.
- 38. (Previously presented) The recombinant
  orthopoxvirus defined in claim 37 wherein the orthopoxvirus is
  selected from the group consisting of a modified vaccinia Ankara
  virus, vaccinia virus Western Reserve, and vaccinia virus
  Copenhagen.
- 39. (Previously presented) The recombinant
  orthopoxvirus defined in claim 37 wherein the orthopoxvirus is the
  modified vaccinia Ankara virus.
  - 40. (Previously presented) The recombinant orthopoxvirus defined in claim 37 wherein the heterologous sequence

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- integrated into the orthopoxvirus in its ATI region is from the ATI region of modified vaccinia Ankara virus.
- 1 41. (Previously presented) The recombinant
  2 orthopoxvirus defined in claim 40 wherein the orthopoxvirus is
  3 selected from the group consisting of a modified vaccinia Ankara
  4 virus, vaccinia virus Western Reserve, and vaccinia virus
  5 Copenhagen.
- 1 42. (Previously presented) The recombinant
  2 orthopoxvirus defined in claim 40 wherein the orthopoxvirus is the
  3 modified vaccinia Ankara virus.
  - 43. (Currently amended) A recombinant orthopoxvirus comprising an ATI region including within an open reading frame of or within the ECORI site of the ATI region an integrated heterologous sequence wherein said recombinant orthopoxvirus is obtained by a method comprising the steps of:
    - (a) transducing a host cell with a vector which comprises an isolated nucleic acid sequence from the ATI region of modified vaccinia Ankara virus that includes at least one restriction enzyme recognition site as an insertion site for the heterologous sequence, that hybridizes under stringent conditions to the nucleic acid sequence of SEQ ID NO:1 or its complementary strand, and that is said nucleic acid sequence capable of integration of the heterologous sequence through homologous recombination into an open

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- reading frame or the ECORI site of the ATI region of an
  orthopoxvirus without interfering with its viral propagation or
  replication efficiency, and at least one heterologous sequence
  inserted within the insertion site into an open reading frame or
  the ECORI site of the isolated nucleic acid sequence;
- (b) infecting said host cell with an orthopoxvirus having an ATI region;
  - (c) integrating the heterologous sequence into an open reading frame or the ECORI site of the ATI region of the orthopoxvirus by homologous recombination between the nucleic acid sequence and a corresponding genomic sequence of the orthopoxvirus to obtain a recombinant orthopoxvirus; and
    - (d) isolating said recombinant orthopoxvirus.
  - 44. (Currently amended) A recombinant orthopoxvirus comprising an ATI region including within an open reading frame of or within the ECORI site of the ATI region an integrated heterologous sequence wherein said recombinant orthopoxvirus is obtained by a method comprising the steps of:
    - (a) transducing a host cell with a vector which comprises an isolated fragment of a nucleic acid sequence from the ATI region of modified vaccinia Ankara virus consisting essentially of at least 200 base pairs of the nucleic acid sequence that is SEQ ID NO:1, that is at least 70% homologous to SEQ ID NO:1 and that includes at least one restriction enzyme recognition site as an insertion site for the heterologous sequence, said isolated

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- fragment of the nucleic acid sequence capable of integration of the 13 heterologous sequence through homologous recombination into an open 14 reading frame or an ECORI site of the ATI region of an 15 orthopoxvirus without interfering with its viral propagation or 16 replication efficiency and at least one heterologous sequence 17 inserted within the insertion site into an open reading frame or 18 the ECORI site of the isolated nucleic acid sequence; 19
- (b) infecting said host cell with an orthopoxvirus having 20 an ATI region; 21
  - (c) integrating the heterologous sequence into an open reading frame or the ECORI site of the ATI region of the orthopoxvirus by homologous recombination between the isolated fragment of the nucleic acid sequence and a corresponding genomic sequence of the orthopoxvirus to obtain a recombinant orthopoxvirus; and
    - (d) isolating said recombinant orthopoxvirus.
- 45. (Currently amended) A recombinant orthopoxvirus comprising an ATI region including within an open reading frame of 2 or within the ECORI site of the ATI region [[a]] an integrated heterologous sequence wherein said recombinant orthopoxvirus is obtained by a method comprising the steps of:
  - (a) transducing a host cell with a vector which comprises an isolated nucleic acid sequence according to SEQ ID NO:1 or its complementary strand from the ATI region of modified vaccinia Ankara virus <del>that includes at least one restriction enzyme</del>

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recognition site as an insertion site for the heterologous 10 sequence, and that is capable of integration of the heterologous 11 sequence into an open reading frame or an ECORI site of the ATI 12 region of an orthopoxvirus without interfering with its viral 13 propagation or replication efficiency, and at least one 14 heterologous sequence inserted within the insertion site into an 15 open reading frame or the ECORI site of the isolated nucleic acid 16 sequence; 17

- (b) infecting said host cell with an orthopoxvirus having an ATI region;
  - (c) integrating the heterologous sequence into an open reading frame or the ECORI site of the ATI region of the orthopoxvirus by homologous recombination between the nucleic acid sequence and a corresponding genomic sequence of the orthopoxvirus to obtain a recombinant orthopoxvirus; and
    - (d) isolating said recombinant orthopoxvirus.
  - 46. (Currently amended) A method of integrating a heterologous sequence into an open reading frame or the ECORI site of the ATI region of an orthopoxvirus to obtain a recombinant orthopoxvirus which comprises the steps of:
  - (a) transducing a host cell with a vector comprising an isolated nucleic acid sequence from the ATI region of modified vaccinia Ankara virus that includes at least one restriction enzyme recognition site as an insertion site for the heterologous sequence, that hybridizes under stringent conditions to the nucleic

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- acid sequence of SEQ ID NO:1 or its complementary strand, and that

  is said nucleic acid sequence capable of integration of the
- heterologous sequence through homologous recombination into an open
- reading frame or the ECORI site of the ATI region of an
- orthopoxvirus without interfering with its viral propagation and
- replication efficiency, and at least one heterologous sequence
- inserted within the insertion site into an open reading frame or
- the ECORI site of the isolated nucleic acid sequence;
- (b) infecting said host cell with an orthopoxvirus having an ATI region;
  - (c) integrating the heterologous sequence into an open reading frame or the ECORI site of the ATI region of the orthopoxvirus by homologous recombination between the nucleic acid sequence and a corresponding genomic sequence of the orthopoxvirus to obtain a recombinant orthopoxvirus; and
    - (d) isolating said recombinant orthopoxvirus.
- 1 47. (Currently amended) The method of integrating a
  2 heterologous sequence into the open reading frame or the ECORI site
  3 of the ATI region of the orthopoxvirus defined in claim 46 wherein
  4 according to step (b) the orthopoxvirus is modified vaccinia Ankara
  5 virus.
  - 48. (Currently amended) A method of integrating a heterologous sequence into an open reading frame or the ECORI site

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- of the ATI region of an orthopoxvirus to obtain a recombinant orthopoxvirus which comprises the steps of:
- (a) transducing a host cell with a vector comprising an isolated fragment of a nucleic acid sequence from the ATI region of modified vaccinia Ankara virus consisting essentially of at least 200 base pairs of the nucleic acid sequence that is SEQ ID NO:1, that is at least 70% homologous to SEQ ID NO:1 and that includes at least one restriction enzyme recognition site as an insertion site 10 for the heterologous sequence, said isolated fragment of the 11 nucleic acid sequence capable of integration of the heterologous 12 sequence into the ATI region of an orthopoxvirus through homologous 13 recombination into an open reading frame or an ECORI site of the 14 ATI region without interfering with its viral propagation or 15 replication efficiency and at least one heterologous sequence 16 inserted within the insertion site; 17
  - (b) infecting said host cell with an orthopoxvirus having an ATI region;
  - (c) integrating the heterologous sequence into an open reading frame or the ECORI site of the ATI region of the orthopoxvirus by homologous recombination between the isolated fragment of the nucleic acid sequence and a corresponding genomic sequence of the orthopoxvirus to obtain a recombinant orthopoxvirus; and
    - (d) isolating said recombinant orthopoxvirus.

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- 1 49. (Currently amended) The method of integrating a
  2 heterologous sequence into an open reading frame or the ECORI site
  3 of the ATI region of the orthopoxvirus defined in claim 48 wherein
  4 according to step (b) the orthopoxvirus is modified vaccinia Ankara
  5 virus.
- 50. (Previously presented) A target cell comprising the recombinant orthopoxvirus defined in claim 37.
- for effecting an immune response against an infectious disease or a proliferative disorder which consists essentially of a therapeutically effective amount of the recombinant orthopoxvirus as defined in claim 37 and in a form capable of producing an immune response against an infectious disease or a proliferative disorder in combination with a pharmaceutically acceptable inert carrier or diluent.
- 52. (Previously presented) A method of effecting an immune response against an infectious disease or a proliferative disorder in an animal subject which comprises the step of administering to said subject a therapeutically effective amount of the pharmaceutical composition defined in claim 51.
- 53. (New) A vector for integration of a heterologous sequence into an open reading frame of or into the ECORI site of

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the ATI region of an orthopoxviral genome having an ATI region, said vector including an isolated nucleic acid sequence from the ATI region of modified vaccinia Ankara virus, that hybridizes under 5 stringent conditions to the nucleic acid sequence of SEQ ID NO:1 or 6 its complementary strand, which includes at least one heterologous 7 sequence inserted into an open reading frame or the ECORI site of 8 the isolated nucleic acid sequence, said at least one heterologous 9 sequence functionally associated with a transcriptional control 10 element thereof, said isolated nucleic acid sequence capable of 11 integration of the heterologous sequence through homologous 12 recombination into an open reading frame or the ECORI site of the 13 ATI region of an orthopoxvirus without interfering with its viral 14 propagation or replication efficiency.

A vector for integration of a heterologous 54. sequence into an open reading frame of or into the ECORI site of the ATI region of an orthopoxviral genome having an ATI region, said vector including an isolated fragment of a nucleic acid sequence from the ATI region of modified vaccinia Ankara virus consisting essentially of at least 200 base pairs of the nucleic acid sequence that is SEQ ID NO:1, which includes at least one heterologous sequence inserted into an open reading frame or the ECORI site of the isolated fragment of a nucleic acid sequence, and which is functionally associated with a transcriptional control element thereof, said isolated fragment of the nucleic acid sequence capable of integration of the heterologous sequence

- through homologous recombination into an open reading frame or an
- 14 ECORI site of the ATI region of an orthopoxvirus without
- interfering with its viral propagation or replication efficiency.